DOPAMINE- dopamine hydrochloride liquid Deseret Biologicals, Inc.

Disclaimer: This homeopathic product has not been evaluated by the Food and Drug Administration for safety or efficacy. FDA is not aware of scientific evidence to support homeopathy as effective.

Drug Facts:

ACTIVE INGREDIENTS:

Dopamine Hydrochloride 8X, 12X, 30X, 200X, 12C, 30C, 60C, 200C.

HOMEOPATHIC INDICATIONS:

For temporary relief of symptoms including headache, reading difficulty, and poor memory.**

**These statements are based upon homeopathic principles. They have not been reviewed by the Food and Drug Administration.

WARNINGS:

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center right away.

If pregnant or breast-feeding, ask a health professional before use.

Tamper seal: "Sealed for Your Protection." Do not use if seal is broken or missing.

KEEP OUT OF REACH OF CHILDREN:

Keep out of reach of children. In case of overdose, contact a physician or Poison Control Center right away.

DIRECTIONS:

1-10 drops under the tongue, 3 times a day or as directed by a health professional. Consult a physician for use in children under 12 years of age.

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INACTIVE INGREDIENTS:

Demineralized Water, 25% Ethanol

QUESTIONS:

Dist. By: Deseret Biologicals, Inc.

469 W. Parkland Drive

PACKAGE LABEL DISPLAY:

DESBIO

NDC 43742-1281-1

HOMEOPATHIC

DOPAMINE

1 FL OZ (30 ml)

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LOT:

Dist. By: Deseret Biologicals, Inc. 469 W. Parkland Drive Sandy, UT 84070 www.desbio.com



HOMEOPATHIC





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DOPAMINE

dopamine hydrochloride liquid

Product Information					
Product Type	HUMAN OTC DRUG	Item Code (Source)	NDC:43742-1281		
Route of Administration	ORAL				

Active Ingredient/Active Moiety					
Ingredient Name	Basis of Strength	Strength			
DOPAMINE HYDRO CHLO RIDE (UNII: 7L3E358N9L) (DOPAMINE - UNII:VTD58H1Z2X)	DOPAMINE HYDROCHLORIDE	8 [hp_X] in 1 mL			

Inactive Ingredients				
Ingredient Name	Strength			
WATER (UNII: 059QF0KO0R)				
ALCOHOL (UNII: 3K9958V90M)				

Packaging					
# Item Code	Package Description	Marketing Start Date	Marketing End Date		
NDC:43742- 1281-1	30 mL in 1 BOTTLE, DROPPER; Type 0: Not a Combination Product	07/27/2018			
Marketing Information					
Marketing Cate	gory Application Number or Monograph Citation	Marketing Start Date	Marketing End Date		
unapproved homeo	athic	07/27/2018			

Labeler - Deseret Biologicals, Inc. (940741853)

Registrant - Apotheca Company, Inc. (844330915)

Establishment			
Name	Address		Business Operations
Apotheca Company, Inc.		844330915	manufacture(43742-1281), api manufacture(43742-1281), label(43742-1281), pack(43742-1281)

Revised: 7/2018 Deseret Biologicals, Inc.